STIC-Biotech/ChemLib

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Chan, Christina

Sent: To: Wednesday, July 27, 2005 4:01 PM Basi, Nirmal; STIC-Biotech/ChemLib

Subject:

RE: Rush search for App. # 10/016,496

Please rush. Thanks Chris

Chris Chan SPE, 1644 TC 1600 New Hire Training Coordinator 571-272-0841 Remsen 3E89

----Original Message----

From: Basi, Nirmal

Sent: Wednesday, July 27, 2005 3:59 PM

To: Char

Chan, Christina

Subject:

Rush search for App. # 10/016,496

Christina I am seeking approval for a RUSH sequence search, as indicated below. If

approved, could you

please forward the search to STIC and cc a copy to me.

Examiner: Nirmal S. Basi

Art Unit 1646

Office: Remsen Building, Room 4D68 Mail Room: Remsen Building, room 4C70

Sequence search:

App. #: 10/016,496 Result format: Paper.

Title: POLYCATION-SENSING RECEPTOR IN AQUATIC SPECIES AND METHODS ____ OF USE THEREOF

Inventors: William Harris et al

Priority Date: 3/27/96 Please search: i) SEQ ID NOs:1 and 2

Search issued, commercial databases and pending databases.

Thanks, Nirmal S. Basi

*******	*******	***********
STAFF USE ONLY	Type of Search	Vendors and cost where applicable
Searcher: Searcher Phone: 2- Date Searcher Picked up: Date Completed: Searcher Prep/Rev. Time: Online Time:	NA#: AA#: Interference: SPDI: S/L: Oligomer: Encode/Transl: Text: Inventor: Litigation:	STN: DIALOG: QUESTEL/ORBIT: LEXIS/NEXIS: SEQUENCE SYSTEM: WWW/Internet: Other(Specify):

Extracellular calcium-sensing receptor precursor (CaSR) (Parathyroid Cell calcium-sensing receptor).
Name=CASR; Syncoyms=GPRC2A, PCARI;

Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Butheria; Primatés; Catarrhini; Hominidae; Homo.

Homo sapiens (Human).

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MAN STANDARD; PRT; 1078 AA.
Q13912; Q16108; Q16109; Q16110; Q16379;
1995 (Rel. 31; Created)
-1995 (Rel. 32; Last sequence update)
-2004 (Rel. 45; Last annotation update)
                                                                                                                                                                                                                                                                                                                      VARIANTS FHH MET-62; CYS-66; MET-138; GLU-143 AND GLN-227.
MEDLINE-95243222; PubMede-7726161;
Chou Y.-H.W., Pollak M.R., Brandl M.L., Toss G., Arnqvist H.,
Atkinson A.B., Papapoulos S.B., Marx S., Brown E.M., Seidman J.G.,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        MEDLINB-95179179, PubMed-7874174;
POSILAK M.R., Brown E.M., Estep H.L., McLaine P.N., Kifor O., Park J., Follak M.R., Brown E.M., Beldman J.G.;
Hebbert S.C., Seidman C.E., Seidman J.G.;
"Autosomal dominant hypocalcaemia caused by a Ca(2+)-sensing receptor
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 VARIANTS FHH GIN-185; LYS-297 AND TRP-795.
MEDLINE=94094324; PubMed=7916660;
POLLAK M.R., Brown B.M., Chou Y.-H.W., Hebart S.C., Marx S.J.,
Steinmann B., Levi T., Seidman C.B., Seidman J.G.;
"Mutations in the human Ca(2+)-sensing receptor gene cause familial hypocalciuric hypercalcemia and neonatal severe hyperparathyroidism.";
Cell 75:1297-1303(1993).
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                SEQUENCE FROM N.A.
Pearce S.H.S., Thakker R.V.;
Submitted (DEC-1994) to the EMBL/GenBank/DDBJ databases
       SEQUENCE OF 1-61 FROM N.A.,
                                                                                                                                                                               "Mutations in the human Ca(2+)-sensing-receptor gene that cause familial hypocalciuric hypercalcemia."; Am. J. Hum. Genet. 56:1075-1079(1995).
                                                                                                                                                                                                                                                                                             Seidman C.B.,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   VARIANT ADH ALA-127.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              "Molecular cloning and functional expression of human parathyroid calcium receptor cDNAs.",
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  TISSUE-Parathyroid;
MEDLINE-95279439; PubMed=7759551; DOI=10.1074/jbc.270.21.12919;
Garrart J.R. Capuano I.V., Hammerland L.G., Hung B.C., Brown B.M.,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Garrett J.E.,
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EDLINE-96343808; PubMed-8756555; DOI=10.1210/en.137.9.3842;
Fraichel M., Zink-Lorenz A., Holloschi A., Hafner M., Flocker
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               uman kidney."
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Hebert S.C., Nemeth E.F., Fuller F.;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   ikle D.D., Ratnam A., Mauro T., Harris J., Pillai S.; Changes in calcium responsiveness and handling during keratinocyte differentiation. Potential role of the calcium receptor."; Clin. Invest. 97:1085-1093(1996).
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  EDLINE-95408281; PubMed=7677761;
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Molecular cloning of a putative Ca(2+)-sensing receptor cDNA from
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              'ISSUE-Kidney;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          EQUENCE OF 643-908 FROM N.A.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Genet. 8:303-307(1994).
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           mutation."
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                                                                                                                     AND VARIANT FHH ALA-39
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CASR HUMAN P41180, Q13 01-PEB-1995

DISEASE: Defects in

CASR are the cause of autosomal dominant

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hypocalciuric hypercalcemia.";
Endocrine 15:277-282(2001).
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               MEDLINE=99415602; PubMed=10487661; DOI=10.1210/jc.84.9.3036; Stock J.L., Brown R.S., Baron J., Coderre J.A., Mancilla E., De Luca F., Ray K., Mericq M.V.; Pautosomal dominant hypoparathyroidism associated with short and premature osteoarthritis.";
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MEDLINE=97442275; PubMed=9298824;
DOI=10.1002/(SICI)1098-1004(1997)10:3<233::AID-HUMU9>3.3.
Ward B.K., Stuckey B.G.A., Gutteridge D.H., Laing N.G., F
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 MEDLINE=96311554; PubMed=87331
Baron J., Winer K.K., Yanoveki
Zimmerman D., Cutler G.B. Jr.;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 MEDLINE=96292293; PubMed=8675635; Pearce S.H.S., Trump D., Wooding C., Grant D.B., Heath D.A., Hughes I.A.,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        "Mutations in the Ca(2+)-sensing receptor gene cause autosomal dominant and sporadic hypoparathyroidism."; Hum. Mol. Genet. 5:601-606(1996).
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              VARIANTS FIH THR-116; HIS-681 AND SER-806, AND VARIANT SER-851.
MEDLINE=96311554; PubMed=8733126; DOI=10.1093/hmg/5.5.601;
Baron J., Winer K.K., Yanoveki J.A., Cunningham A.W., Laue L.,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            "Calcium-sensing receptor mutations in and neonatal hyperparathyroidism."; J. Clin. Invest. 96:2683-2692(1995).
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            VARIANT FIH VAL-616.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            "A novel mutation (L174R) in the Ca2+-sensing with familial hypocalciuric hypercalcemia.";
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Ratajczak
         nyperpassa. ... the homozygous form of FHH.
DISEASE: Defects in CASR are the cause of
MIN:601198]; in which
                                                                                                                                                                                                                                         TISSUE SPECIFICITY: Found in kidney, but not in brain, lung, liver, heart, skeletal muscle, or placenta.

DISEASE: Defects in CASR are the cause of familial hypocalciuric hypercalcemia, type 1 (FHH) [MIM:145980]; in which the receptor hypercalcemia, type 1 (FHH) [MIM:145980]; in which the receptor hypercalcemia, type 1 (FHH) [MIM:145980]; in which the receptor
                                                                                                                                                                                                                                                                                                                                                                                                                    messenger system.
SUBCELLULAR LOCATION: Integral membrane
ALTERNATIVE PRODUCTS:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                       FUNCTION: Senses changes in the extracellular concentration calcium ions. The activity of this receptor is mediated by a protein that activates a phosphatidylinositol-calcium second
                                                                                   DISEASE: Defects in CASR are the cause of neonatal severe primary hyperparathyroidism (NSHPT) [MIM:239200]; in which the receptor has reduced activity. NSHPT is a rare autosomal recessive lifethreatening disorder characterized by very high serum calcium concentrations, skeletal demineralization, and parathyroid
                                                                      hyperplasia. In some instances
                                                                                                                                                                                             hypercalcemia, relative hypocalciuria, and inappropriately normal
                                                                                                                                                                                                                has reduced activity. FHH is characterized by altered ca
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              familial hypocalciuric h
Mutat. 10:233-235(1997).
                                                                                                                                                                                                                                                                                                                                                               IsoId=P41180-1; Sequence=Displayed;
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                                                                                                                                                                                levels.
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Metab. 80:2594-2598(1995)
                                                                                                                                                                                                                                                                                                                            Sequence=VSP
                                                                                                                                                                                                                                                                                                                                                                                                   splicing, Named isoforms=2;
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Pound in kidney, but ...

Pound in kidney, but ...

Pound in kidney, but ...
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                                                                  has been demonstrated
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             autosomal dominant the receptor is
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Matches 782
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EMBL; U20759;
EMBL; U50760;
EMBL; D50855;
EMBL; S83176;
EMBL; S83176;
EMBL; S68032;
EMBL; S68033;
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PROSITE; PS00979; G_PROTEIN_RECEP_F3_1;
PROSITE; PS00980; G_PROTEIN_RECEP_F3_2;
PROSITE; PS00981; G_PROTEIN_RECEP_F3_3;
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Pfam; PF01094; ANF receptor; 1.
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InterPro, IPR000337; GPO
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                   GO:0005887; C:integral to plasma membrane; TAS.
GO:0004930; F:G-protein coupled receptor activity; TAS.
GO:0004930; P:protein coupled receptor activity; TAS.
GO:0006474; P:calcium ion homeostasis; TAS.
GO:0005513; P:calcium ion sensing; TAS.
GO:0007585; P:chemosensory behavior; TAS.
GO:0007185; P:crotein coupled receptor protein signalin.
GO:0007853; P:morphogenesis; TAS.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                      GO:0001503;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  parathyroid hormone. Symptoms are seizures, tetany and cramps. SIMILARITY: Belongs to the G-protein coupled receptor 3 family.
                            241
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             hypoparathyroidism (FIH) [MIM:146200]. FIH is characterized by hypocalcemia and hyperphosphatemia due to inadequate secretion
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; $81755;
; P23385;
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                                                                                                                                                                                                                                                                                      Similarity.
                                                                                                                                                                                                                                         MAQLHCQLLFLGFTLLQSYNVSGYGPNQRAQKKGDIILGGLFFIHFGVAAKDQDLKSRPE
               IDFSEMISQYYTQKQLEFIADVIQNSSAKVIVVFSNGFDLEFLIQEIVRRNITDRIWLAS
                                                                  KNEYKAPIRTIPNDEQQATAMABIIEHFQWNWVGTLAADDDYGRPGIDKFREBAVKRDIC
                                                                                                                        QNKIDSLANLDEFCNCSDHIPSTIAVVGATGSGISTAVAANLLGLFYIPQVSYASSSRLLSN
IDFSELISQYSDEEEIQHVVEVIQNSTAKVIVVFSSGPDLEPLIKEIVRRNITGKIWLAS
                                                     KNOFKSFLRTIPNDEHQATAMADIIEYFRWNWVGTIAADDDYGRPGIEKFREEAEERDIC
                                                                                                          QNKIDSLNIDEFCNCSEHIPSTIAVVGATGSGVSTAVANLLGIFYIPQVSYASSSRLLSN
                                                                                                                                                             SVECIRYNPRGPRWLOAMIPAIBBINSSPALLPNLTLGYRIPDTCNTV8KALBATLSFVA
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AAB46873.
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RESULT 3
CASR_BOVIN
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                                                                                                                                                                               Cell calcium-sensing receptor).
Name=CASR; Synonyms=GPRC2A, PCAR1;
Bos taurus (Bovine).
                                                                                                                                                                                                                                                                                                                                                 CASR BOVIN
                                                                                                                                    Eukaryota; Metazoa;
Mammalia; Eutheria;
TISSUE-Parathyroid;
MEDLINE-94077182; PubMed-8255296; DOI-10.1038/366575a0;
                                           SEQUENCE FROM N.A.
                                                                                 NCBI_TaxID=9913;
                                                                                                                 Bovinae;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - between Bioinformatics Institute. Three are no restrictions on its the European Bioinformatics Institutions as long as its content is in no way use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (see http://www.isb-sib.ch/announce/
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         -I- FUNCTION: Senses changes in the extracellular concentration calcium ions. The activity of this receptor is mediated by a protein that activates a phosphatidylinositol-calcium second
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SIGNAL
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Pfam; PF01094; ANP_receptor; 1.

PRINTS; PR00248; GPCRMGR.

PROSITE; PS00979; G PROTEIN RECEP F3 1; 1.

PROSITE; PS00980; G PROTEIN RECEP F3 2; 1.

PROSITE; PS00981; G PROTEIN RECEP F3 3; 1.

PROSITE; PS50259; G PROTEIN RECEP F3 4; 1.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             entities requires a license agreement (S or send an email to license@isb-sib.ch)
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PIR; S40476; S40476.
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Interpro; IPR000068; Ca_sens_receptor.
Interpro; IPR000337; GPCR_Mgr.
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SUBCELLULAR LOCATION: Integral membrane protein.
SIMILARITY: Belongs to the G-protein coupled receptor 3 family.
              61
                                                                                              Similarity
ATKCIRYNERGERWLOAMIEAIEEINNSWTELPNITLGYRIEDTCNTVSKALEATLSFVA 120
                                      maqlhcqllfigftllosynvsgygdnoraqkkgdiilgglffihfgyaakdodlksrpb
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                                                                                    Conservative
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5 (Potential):
Cytoplasmic (Potential).
6 (Potential).
Extracellular (Potential).
7 (Potential).
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2 (Potential)
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Pred. No. 4e-271;
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SVECIRYNFRGFRWLQAMIFAIBEINSSPALLPNMTLGYRIFDTCNTVSKALBATLSFVA

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GLTMEMQ-----RCSTQKVSFGSGTVTLSLSFEETGRYATLSRTARSRNSADGRSGDD
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PROSITE; PS00979; GENOTEIN RECEP F3 2; 1.
PROSITE; PS00980; GENOTEIN RECEP F3 3; 1.
PROSITE; PS00981; GENOTEIN RECEP F3 3; 1.
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Pfam; PF01094; ANF receptor; 1.
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RGD; 2277; Casr.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            to nerve terminals.";

Proc. Natl. Acad. Sci. U.S.A. 92:3161-3165(1995).

I- FUNCTION: Senses changes in the extracellular concentration calcium ions. The activity of this receptor is mediated by a protein that activates a phosphatidylinositol-calcium second
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STRAIN=Wistar;
WEDLINE=95241465; PubMed=7724534;
Ruat M., Snowman A.M., Snyder S.H.;
"Calcium sensing receptor: molecular cloning
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      STRAIN=Sprague-Dawley; TISSUE=Kidney outer medulla; MEDLINE=95116508; PubMed=7816802; Riccardi D., Park J., Lee W., Gamba G., Brown E.M., "Cloning and functional expression of a rat kidney e calcium/polyvalent cation-sensing receptor."; Proc. Natl. Acad. Sci. U.S.A. 92:131-135(1995).
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Mammalia; Butheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
NCBI_TaxID=10116;
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4 (Potential).

Extracellular (Potential).

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Cytoplasmic (Potential).

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Extracellular calcium-sensing receptor.

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                                                                                                                                                                                                                                                                                                                                                                     Query Match
Best Local Similarity
Matches 778; Conserv
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       InterPro; IPR001828; ANF receptor.
InterPro; IPR00068; Ca sens receptor.
InterPro; IPR0100337; GPCR MgT.
InterPro; IPR011500; WCD3G GPCR.
Pfam; PF00003; 7tm 3; 1.
Pfam; PF01094; ANF receptor; 1.
Pfam; PF07562; WCD3G; CASENSINGR.
PRINTS; PR00248; GPCRMGR.
PRINTS; PR00248; GPCRMGR.
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Q1-JUN-2003 (TYEMBLrel. 24, Cre
O1-JUN-2003 (TYEMBLrel. 25, Las
O1-CCT-2003 (TYEMBLrel. 25, Las
Calcium sensing receptor.
Raftus morvegicus (Rat).
Eukaryota; Metazoa; Chordata; C:
Mammalia; Butheria; Rodentia; S:
NCBI TaxID-10116;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         PROSITE; PS00979; G PROTEIN RECEP F3 1;
PROSITE; PS00980; G PROTEIN RECEP F3 2;
PROSITE; PS00981; G PROTEIN RECEP F3 3;
PROSITE; PS50259; G PROTEIN RECEP F3 4;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  GO; GO:0016020; C:membrane; IEA.
GO; GO:0008067; F:metabotropic glutamate,
GO; GO:0004872; F:receptor activity; IEA.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Bukoski R. Jain K., Wang Y.;
Submitted (JAN-2003) to the EMBL/GenBank/DDBJ
EMBL; AY214122; AA059490.1; -.
HSSP; P23385; 1EWK.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              STRAIN=Wistar; TISSUE=Dorsal root Bukoski R., Bian K., Wang Y.;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Q80ZA8
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MAQLHCQLLFLGFTLLQSYNVSGYGPNQRAQKXGDIILGGLFFIHFGVAAKDQDLKSRPE
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73.5%;
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Last sequence update)
Last annotation update)
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Pred. No. 3.5e-270;
37; Mismatches 140;
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Sciurognathi; Muridae;
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Murinae; Rat
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